

MICRO-INVASIVE BREAST BIOPSY DEVICE

Related Application

This application claims the benefit of U.S. provisional applications Serial No. 60/278,128, filed March 23, 2001 and Serial No. 60/281,848, filed April 5, 5 2001 and Serial No. 60/305,178 filed July 13, 2001 and Serial No. 60/322,909, filed September 17, 2001 and Serial No. 60/342,436, filed December 21, 2001, the disclosure of each of which is incorporated in its entirety herein by reference.

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Field of the Invention

The present invention relates generally to medical devices, and, more particularly, to micro-invasive devices 15 and methods for removing breast tissue for biopsy and treatment.

Background of the Invention

The medical industry is constantly evolving through the adaptation of improved pharmaceutical, biotechnology, 20 and medical device products and procedures. Techniques and technologies are being developed to treat internal areas of the body through less invasive means.

It is often desirable and frequently necessary to remove a portion of tissue from humans and other animals, 25 particularly in the diagnosis and treatment of patients with cancerous tumors, pre-malignant conditions and other diseases or disorders. Typically, in the case of cancer, when the physician establishes by means of procedures such as palpation, x-ray or ultrasound imaging that suspicious 30 circumstances exist, a biopsy is performed to determine whether the cells are cancerous. Biopsy may be done by an

open or percutaneous technique. Open biopsy removes the entire mass (excisional biopsy) or a part of the mass (incisional biopsy). Percutaneous biopsy, on the other hand, is usually done with a needle-like instrument and may be either a fine needle aspiration (FNA) or a core biopsy. In FNA biopsy, individual cells or clusters of cells are obtained for cytologic examination and may be prepared such as in a Papanicolaou smear. In core biopsy, as the term suggests, a core or fragment of tissue is obtained for histologic examination which may be done via a frozen section or paraffin section. The type of biopsy utilized depends in large part on circumstances present with respect to the patient and no single procedure is ideal for all cases. However, core biopsy is extremely useful in a number of conditions and continues to be used frequently by the medical profession.

To arrive at a definitive tissue diagnosis, intact tissue is needed from an organ or lesion within the body. In most instances, only part of the organ or lesion need be sampled. However, the portions of tissue obtained must be representative of the organ or lesion as a whole. In the past, to obtain tissue from organs or lesions within the body, surgery had to be performed to locate, identify and remove the tissue. With the advent of medical imaging equipment (x-rays and fluoroscopy, computed tomography, ultrasound, nuclear medicine, and magnetic resonance imaging) it has become possible to identify small abnormalities even deep within the body. However, definitive tissue characterization still requires obtaining adequate tissue samples to characterize the histology of the organ or lesion. For example, mammography can identify non-palpable (not perceptible by touch) breast abnormalities earlier than they can be diagnosed by physical examination. Most non-palpable breast abnormalities are benign; some of them are malignant. When

breast cancer is diagnosed before it becomes palpable, breast cancer mortality can be reduced. However, it is often difficult to determine if pre-palpable breast abnormalities are malignant, as some benign lesions have
5 mammographic features which mimic malignant lesions and some malignant lesions have mammographic features which mimic benign lesions. Thus, mammography has its limitations. To reach a definitive diagnosis, tissue from within the breast must be removed and examined under a
10 microscope. Early biopsy techniques for reaching a definitive tissue diagnosis for non-palpable breast disease required a mammographically guided localization, either with a wire device, visible dye, or carbon particles, followed by an open, surgical biopsy utilizing one of these
15 guidance methods to lead the surgeon to the non-palpable lesion within the breast.

Open surgical breast biopsies have many drawbacks. They can be disfiguring, expensive and are imperfect. Open surgical biopsies also carry a small mortality risk (due to
20 the risks of anesthesia) and a moderate morbidity rate (including bleeding, infection, and fracture or migration of the localizing wire). In cases where multiple lesions are present in the breast, a surgeon is reluctant to biopsy each lesion due to the large tissue mass that must be extracted with each lesion. The most convenient lesion may
25 be taken which most often results in an incomplete diagnosis.

Percutaneous breast biopsy techniques are more desirable in many instances, particularly in light of modern imaging techniques which are able to pinpoint
30 nonpalpable tissue masses in the breast and consequently, the increased frequency of biopsies that are necessary for diagnosis of these tissue masses. A well known instrument used quite extensively for core biopsies in the past is manufactured by Travenol Laboratories of Deerfield, Ill.,
35 and is sold under the mark "TRU-CUT." This manual biopsy

instrument at one time enjoyed as much as 98% of the market for such devices. As disclosed in U.S. Pat. No. 3,477,423, the instrument comprises a two-piece assembly: an outer cutting cannula mounted to one hub member and an inner stylet with a sampling notch ground into it mounted to a second hub, with the hubs being slidably interlocked. The instrument is assembled and placed into the body with the outer cutting cannula just to the rear of a lancet point or beveled distal end of the stylet. Upon inserting the device up to or in front of the area to be biopsied, advancement of the assembly is halted. The stylet is manually advanced distally of the cannula with the cannula held stationery. Upon advancement of the stylet, the specimen notch is exposed. Tissue surrounding the stylet prolapses into the specimen notch and the cutting cannula is then manually advanced distally over the stylet, slowly shearing off the tissue entrapped in the stylet's specimen notch. The instrument is then either (a) withdrawn and the stylet advanced distally to expose the tissue for preparation for study or (b) left in place and only the stylet is proximally removed from within the cannula so a determination of successful sampling may be made. If the sampling was not successful, the stylet may be reinserted into the cannula, which remains positioned within the patient, and an attempt to reposition the assembly of stylet and cannula and repeat sampling can be made.

Such a technique using this basic design of a biopsy instrument is referred to as a manual technique. One drawback to the manual technique is that it requires a great deal of manual dexterity and motor coordination, along with the use of both hands, to advance the stylet while maintaining the position of the cannula and then to maintain the position of the stylet while advancing the cannula. Another drawback is that the cannula is advanced relatively slowly, resulting in an extremely poor cutting

action and allowing the surrounding tissue an opportunity to collapse, thus making no use of the stored kinetic energy in the material being severed. Further disadvantages are encountered when the tissue volume to be sampled 5 contains areas of higher density than that of surrounding tissue, such as areas of calcification commonly associated with certain types of cancerous growths. A manually inserted sampling device is often incapable of penetrating the denser area of tissue which merely deflects the course 10 of the cannula/stylet structure around the dense area and into the more compliant surrounding tissue. In the late 1980's, two different stereotactic guidance systems were modified to allow the guiding portion of each system to accommodate spring powered devices such as the Biopty® 15 (Bard Radiology) gun. As used herein, the term "gun" is used to refer to a tissue sampling device designed for "one-handed" operation. A feature common to many of these devices is the shape of the device being adapted to fit the hand of a medical practitioner with a pistol-like grip, 20 complete with a triggering mechanism. Free-hand ultrasound guidance techniques were also developed to guide the Biopty® gun to breast lesions detected by ultrasound.

With image-guided percutaneous core breast biopsy, it should be possible to greatly reduce the number of open, 25 surgical breast biopsies performed. However, there are limiting factors with image-guided breast biopsies. Conventional, manually operated two-step devices are awkward to manipulate, and the tissue samples obtained thereby are often unsatisfactory.

30 A variety of automatic and semiautomatic biopsy instruments have been developed which are spring loaded gun-type devices. A biopsy gun currently used is described in U.S. Pat. No. 34,056, entitled "TISSUE SAMPLING DEVICE," issued to Lindgren et al. Additional examples of 35 biopsy gun devices are disclosed in U.S. Pat. Nos.

4,600,014 and 4,958,625.

Such devices use a design comprising a handle held in a physician's palm, and a guide tube extending forwardly of the handle. A cannula is slidably disposed within the guide tube and is movable from within the guide tube forwardly out of the distal end of the guide tube. A sampling stylet is telescopically disposed within the cannula and projects from the rear of the handle. In an automatic mode of operation, the cannula, when in the retracted position, is spring loaded by means of a compressed spring. A release lever, which works against the compressed spring, is activated to release compression of the spring which then expands and pushes the cannula outwardly over the stylet. This instrument, as stated, requires two handed operation. Also, since the stylet is not removable proximally from within the handle, the entire instrument must be withdrawn to obtain access to the sample.

A fully automatic instrument manufactured by Radioplast, Inc. of Sweden is described in U.S. Pat. No. 4,699,154. This instrument comprises a reusable, spring-loaded box-shaped housing or handpiece, which activates a disposable cannula and stylet set. Both the stylet and cannula are activated in rapid succession. The instrument has the advantage of reducing the dexterity and motor coordination necessary in the use of manual devices and also eliminates the slow cutting action of the manually advanced cannula, replacing it with a very quick, clean cut. This instrument, however, also has its drawbacks. First, the reusable handpiece is very large, heavy, cumbersome, and expensive. They are also typically spring-powered devices and must be manually cocked with some sort of plunger bar. Such "cocking" of the gun requires considerable force and the gun must be cocked for each biopsy cut. When actuated, the springs provided in the gun accelerate the needles until a mechanical stop position is

reached which can create a loud snapping noise and jerking motion which is a problem both to the physician and the patient. A further drawback is encountered in automatically activating both the stylet and the cannula, as opposed to activating the stylet manually, in that the rapid speed at which the cannula follows the stylet into the tissue does not allow much tissue to collapse into the specimen notch, limiting the size of the sample.

U.S. Pat. No. 5,183,054, entitled "ACTUATED BIOPSY CUTTING NEEDLE WITH REMOVABLE STYLET," issued to Burkholder et al., discloses a biopsy device having a tubular cannula through which a stylet, having a stylet cavity near the distal end, is placed. The stylet is removable from the cannula and removed from the biopsy device through the housing so that the tissue sample obtained by the biopsy device may be manually retrieved while the cannula remains in place within the patient, near the area being sampled. Thereafter, the stylet may be reinserted through the housing and cannula into the patient's tissue where additional tissue samples may be obtained. In this way, trauma to the tissue that ordinarily occurs upon reinsertion of the cannula and stylet is minimized.

U.S. Pat. No. 5,234,000, entitled "AUTOMATIC BIOPSY DEVICE HOUSING A PLURALITY OF STYLETS," issued to Hakky et al. describes a biopsy device for taking a plurality of samples of tissue from a living being. The device comprises a housing having a portion arranged to be held by a person using the device, a cannula having a proximal portion and a distal portion and being coupled to the housing. A plurality of stylets are located in the housing, with each of the stylets having a proximal end, a distal end, and a tissue receiving notch located adjacent the distal end. Each stylet is individually propelled through the cannula into the body so that a portion of the tissue prolapses into the notch.

There currently exists a need for a more effective microsurgical device for obtaining a sample of breast tissue from a patient.

5 Summary of the Invention

New apparatus and methods for removing breast tissue and/or other material from a human breast have been discovered. The present invention provides apparatus, for example, micro-invasive apparatus, to remove tissue or 10 other material from a target area of a breast to provide one or more benefits, such as diagnostic benefits, therapeutic benefits and the like.

The apparatus of the invention are useful for removing unwanted, diseased, or even healthy bodily materials, for 15 example, tissues, for medical treatment and/or therapeutic purposes. Advantageously, the present invention is suitable for use in many surgical settings and is suitable for performing various material removal procedures using methodologies, for example, in terms of methods of 20 introducing the apparatus into the body and removing the apparatus from the body, which are substantially analogous to conventional surgical techniques. Necessary or desirable adaptations of the apparatus of the present invention for specific medical treatment, e.g., diagnostic, 25 and therapeutic purposes will be readily appreciated by those of skill in the art.

Accordingly, apparatus for removing tissue from a target area of a breast are provided. In one broad aspect, the apparatus comprise a handpiece and a tissue removal 30 element connected or coupled thereto. The tissue removal element includes a cannula, for example, a substantially rigid or flexible cannula, and a rotational element disposed in the cannula. The rotational element is connected to a source of rotational energy, for example, a

motor. The rotational element is disposed at least partially in the cannula. The cannula includes an open distal tip structured to be placed in a breast, and preferably a proximal end connected, for example, removably connected, to the handpiece. The tissue removal element is structured and effective to draw breast tissue or other material from the target area or site, for example, into the open distal tip, in response to, for example, as a result of, rotation of the rotational element relative to the cannula.

In one embodiment, the rotational element is structured to at least assist in drawing material from a breast into the cannula. For example, the rotational element and the cannula cooperatively engage to form or create a source of suction sufficient to draw the tissue or other material into the cannula in response to rotation of the rotational element relative to the cannula. Advantageously, the cannula, in particular the interior hollow space formed or defined by the cannula, and the rotational element are sized and positioned, relative to each other, to create a source of suction or pumping action in response to the rotational element rotating relative to the cannula. Without wishing to limit the invention to any particular theory of operation, it is believed that this functioning of the cannula/rotational element combination is at least somewhat analogous to the functioning of a pump, for example, a pump based on the principles of the "Archimedes' screw", causing the material to be drawn or pulled or pumped into the open distal tip of the cannula and through the cannula in being removed from the target area of the human/animal body.

Preferably, the suction/pumping action created or formed by the cannula/rotational element combination is itself sufficient and effective so that no other, for example, no additional or supplemental, source of suction

or pumping action is employed, needed or required to effectively remove material from the target area in accordance with the present invention.

In one embodiment of the invention, the rotational element includes a shaft and one or more outwardly extending projections, for example threads, preferably having a substantially helical configuration. Advantageously, the rotational element includes a distal portion with such projections or threads. The proximal portion of the rotational element may or may not include such projections or threads. In a very useful embodiment, the proximal portion is substantially free of such projections or threads.

The distal portion of the rotational element, in a useful embodiment, extends beyond the open distal tip or inlet of the cannula, for example, by a distance in a range of about 0.02 inches to about 1 inch beyond the open distal tip of the cannula. For example, the distal portion may extend a distance equal to at least about one-half of the spacing between adjacent projections or threads. The rotational element distal portion may extend a distance equal to more than about one spacing, for example, about two spacings or more between adjacent projections or threads beyond the open distal tip of the cannula. More specifically, the rotational element advantageously further includes an elongated shaft having a proximal portion which is substantially smooth to allow sufficient annular space between the shaft and cannula for removal of material.

The cannula may be of any suitable size. However, in order to obtain the reduced invasiveness benefits of the present invention, it is preferred that the cannula size have an outer diameter of no greater than about 5 mm and more preferably about 2.0 mm or smaller.

It has unexpectedly been found that the present apparatus including such small size cannulas not only

provide for reduced, or even micro, invasive procedures (which reduce surgical trauma and promote healing) but also are effective in removing materials from a body to achieve therapeutic benefits, for example, therapeutic removal of healthy or diseased breast tissue, fluidic cystic materials, soft tissue tumors, and the like.

In one embodiment of the invention, the open distal tip of the cannula is angled or is beveled with respect to a longitudinal axis of the cannula. Alternatively, the open distal tip is substantially perpendicular with respect to the longitudinal axis of the cannula.

The present apparatus advantageously includes a tissue collection chamber in communication, for example, fluid communication, with the cannula and structured to collect and contain material, for example breast tissue, that is passed through the cannula. The collection chamber preferably is structured to facilitate quantification and/or other analysis of the removed material. In one particularly useful embodiment, the collection chamber comprises a substantially transparent conical section removably engaged to a housing of the handpiece and preferably circumscribing a portion, for example, the proximal portion of the shaft of the rotational element.

The cannula and/or the rotation element, preferably both, advantageously are structured to be manually deformable, for example, to enable the physician to alter the normal configuration, for example, the normal substantially straight configuration, thereof to create a curved configuration if desired, for example, to deal with the material removal application at hand.

In another broad aspect of the present invention, methods of removing material from a breast of a human or an animal are provided. Such methods comprise placing into a breast of a human or an animal a cannula having an open distal tip and a rotational element disposed at least

5 partially in the cannula and rotating the rotational element relative to the cannula, thereby at least assisting in drawing a material from the breast into the open distal tip of the cannula. The method preferably further comprises passing the material from the breast through the cannula. Apparatus in accordance with the present invention described herein, can be advantageously employed in accordance with the present methods.

10 The cannula used in accordance with the present methods preferably have outer diameters of about 5 mm or less, or about 2.0 mm or less.

15 The placing step of the present methods preferably includes percutaneously introducing the cannula into the human or animal breast, and positioning the open distal tip of the cannula in close proximity to the material to be removed. The cannula and rotational element preferably are sized and positioned relative to each other so that the rotating step is effective in drawing the material from the breast of a human or an animal into the open distal tip of 20 the cannula. Preferably, the material from the breast is removed without applying additional suction or aspiration to the open distal tip of the cannula.

25 In one very useful embodiment, the rotating of the rotational element relative to the cannula is effective to draw the material into the cannula as a substantially single, continuous often times substantially cohesive piece. Thus, although some shearing and/or cutting of the material to be removed may occur in accordance with the present invention, for example, so that the removed 30 material is compatible with the space within the cannula through which the material is to be moved proximally, the present apparatus and methods preferably are not based on cutting or chopping the material to be removed into small discrete segments.

The present methods preferably further comprise collecting the material removed and/or observing and/or otherwise testing the material removed.

Any suitable material can be removed from the body of 5 a human or an animal using the present apparatus and/or methods. Preferably, such material to be removed can be effectively removed using the present apparatus and/or methods without employing or requiring additional suction or aspiration, beyond that formed or created by the 10 rotation of the rotational element relative to the cannula.

Advantageously, the material to be removed is soft and/or semi-solid and/or a viscous flowable material and/or a material which is at least somewhat free to move toward a source of lower pressure or suction.

15 Incorporated herein by this specific reference are U.S. Patent application for Micro-invasive Tissue Removal Device, having Serial No. (attorney docket no. D-3034), filed on even date herewith, and commonly assigned herewith, and U.S. Patent application for Micro-invasive 20 Nucleotomy Device and Method, having Serial No. (attorney docket no. D-3039) filed on even date herewith, and commonly assigned herewith.

Each and every feature described herein, and each and 25 every combination of two or more of such features, is included within the scope of the present invention provided that the features included in such a combination are not mutually inconsistent.

The present invention and the objects and advantages thereof will be more clearly understood and appreciated 30 with respect to the following Detailed Description, when considered in conjunction with the accompanying Drawings.

Brief Description of the Drawings

Fig. 1 shows a side view of a micro-invasive breast tissue removal apparatus in accordance with the present invention including a handpiece and a tissue removal element connected thereto.

Fig. 2 is a partial cross-sectional view of the apparatus taken along line 2-2 of Fig. 1.

Fig. 3 is a partial cross sectional view of a preferred distal tip of the tissue removal element of the apparatus in accordance with the present invention; and

Figs. 4 is side view of the apparatus shown in Fig. 1, with the tissue removal element having a curve for facilitating access to breast tissue.

Figs. 5 and 6 each show a partial cross sectional view of an alternative distal tip of the tissue removal apparatus of the present invention.

Detailed Description

Turning now to Figs. 1 and 2, a micro-invasive tissue removal apparatus for removing tissue or other material from a breast, in accordance with the present invention is shown generally at 10. The apparatus 10 generally comprises a handpiece 14 and a tissue removal mechanism 16 to be described in detail hereinafter.

The handpiece 14 is preferably sized and contoured to fit comfortably within a palm of a surgeon, and includes for example a molded plastic housing 22. As shown in Fig. 2, the housing 22 of the handpiece 14 encloses a small motor 24 and a power supply, for example a 9 volt battery 26 for driving the tissue removal mechanism 16. Suitable electrical connectors 27 are provided. For convenient, one handed operation, an ON/OFF switch 28 is preferably provided on a recessed, lateral portion 29 of the housing 22.

Turning now as well to Fig. 3, the tissue removal mechanism 16 generally includes a cannula 30 and a rotatable element 34 disposed therein. As shown most clearly in Fig. 3, the cannula 30 includes a distal portion 40 defining an inlet 42 for receiving tissue drawn from a target area within a patient. The inlet 42 is defined, for example, by flat, distal edge 44 of the cannula 30. The distal edge 44, in the embodiment shown in Fig. 3, lies along a plane that is substantially perpendicular with respect to the longitudinal axis of the cannula 30. During operation of the apparatus 10, as will be described in greater detail hereinafter, breast tissue and/or other material is drawn, suctioned or pumped, through the inlet 42 and into a cylindrical bore 46 defined between the cannula 30 and a shaft 50 of the rotatable element 34.

In a preferred embodiment of the invention, such as shown in Figs. 1-3, the tissue removal mechanism 16 is structured to draw breast tissue or other material into the cannula 30 by a pumping action produced by rotation of the rotatable element 34, preferably without the use of supplemental aspiration or other means for drawing tissue into the threaded distal portion 52 or cannula 30. In other words, the rotational element 34 and the cannula 30 are designed to cooperatively engage to form a source of suction that is, in itself, sufficient to draw the material into the cannula 30. Advantageously, the present invention has been found to be safe and highly effective for removing soft tissues from a breast of a patient, for example, less invasively, without being connected to external sources of aspiration or other external machines and devices. In the preferred embodiment of the invention, the rotational element 34 includes a distal portion 52 which extends beyond the open distal tip (defined by edge 44) of the cannula 30. More preferably, the distal portion 52 extends a length of about 0.066 inches beyond the

cannula distal edge 44. A blunt, rounded tip 53 of the rotational element 34 is preferably provided.

As shown, the rotational element 34 includes one or more outwardly extending projections, for example threads such as helical threading 56 shown, disposed about at least a portion of the shaft 50, for urging tissue into the bore 46. Preferably, outer radial edge 58 of the threading 56, or other projection, is disposed closely proximate an inner wall 62 of the cannula. As shown, the distal end 52 of the rotational element 34 extends at least one-half thread turn beyond the cannula inlet 42. This structure allows breast tissue or other material within the breast to prolapse between the outer, distal-most threading turns, and be pulled into the inlet without necessarily being discretely cut or severed by the threading 56. The present invention is designed such that upon insertion of the open distal tip of the cannula 30 into the target region of the breast of a body, tissue or other material will prolapse into and at least partially fill the open spaces between the projections or threading 56. Rotation of the rotational element 34, for example at about 12,000 RPM, causes the material to be pulled in a proximal direction proximally into the bore 46, for example, as a continuous piece or strand of material.

Although the threading 56 is only shown as a single thread located on the distal portion 52 of the rotational element 34, it is to be appreciated that in some embodiments of the invention, the threading 56 may involve multiple threads, and/or may be disposed on more proximally located portions of the rotatable element shaft 50. Furthermore, although only about 4.5 turns of threading 56 are shown, it is to be appreciated that in some embodiments of the present invention, fewer or more than 4.5 turns of threading 56 may be provided. It is also contemplated by the present invention that rather than

continuous threading 56, the shaft 50 may be provided with discontinuous threading. It is contemplated that with appropriate modifications and the like, these and other structures may be provided which would operate in a manner similar to the pumping action provided by the structure shown.

Preferably, the cannula 30 has an outer diameter of less than about 5 mm, for example, an outer diameter of about 2.0 mm or less. The cannula 30 is made of any suitable, medical grade material or materials, but is preferably somewhat rigid but bendable.

Advantageously, as will be appreciated by those of skill in the art, the apparatus 10 of the present invention is minimally invasive to the patient. For example, the cannula 30 can be introduced into the target area of the breast of the patient by means of a conventional, rigid stylet (not shown) disposed through the cannula 30 (detached from the handpiece 14). The cannula/ stylet are introduced percutaneously through the skin, underlying muscle/fatty tissue and into the target area within or adjacent a breast such that the inlet 42 is positioned within or closely adjacent the target tissue or other material. The stylet is then removed and the cannula 30 is left in place. The rotational element 34, attached to the handpiece 14, is then introduced into the cannula 30. This procedure may be facilitated through the use of fluoroscopy or x-ray imaging techniques as known in the art, which do not require direct endoscopic or direct viewing of the target tissue.

Advantageously, unlike prior art surgical breast tissue removal devices, the action of the tissue removal mechanism 16 urges tissue into the cannula 30 in many instances a substantially continuous segment rather than in relatively smaller, distinct portions of the tissue. Generally, the cannula 30 and rotational element 34 are

structured to cooperatively function in a manner that will form a source of suction within the cannula 30 when the rotational element 34 is rotated while the cannula inlet 42 is disposed within the target tissue. It has been found
5 that the level of suction so created is sufficient to gently and effectively draw soft tissue, for example gelatinous, viscous, or any suitable tissue that can be drawn by the action of the present invention into the cannula without need for any other, for example,
10 supplemental, source of suction applied to the inlet 42. For example, the suction formed or created is sufficient to pull or soft tissues into the open tip without causing damage to other structures.

The tissue removal mechanism 16 can be left to remain
15 in substantially the same position within the target area of the breast during the tissue removal procedure, or alternatively may be advanced, or withdrawn during the procedure, for example in a direction along the longitudinal axis of the cannula in order to facilitate
20 tissue removal.

Fig. 4 shows another advantageous feature of the present invention. The tissue removal mechanism 16 may be structured to be deformed, for example, manually deformed, into a curve shape such as shown. The flexibility and
25 deformability of the tissue removal mechanism 16 allows custom shaping or curving of the apparatus 10 for further facilitating access to tissue.

Unlike prior art devices designed to remove substantially liquid substances, the present invention can
30 be used to remove highly viscous substances.

Fig. 5 shows an alternative cannula distal portion 40a, which is beveled, includes sharp distal tip 80, and a relatively wider inlet 42a than inlet 42. Also shown is a narrower threading 56a (relative to threading 56 of Fig. 3)
35 on rotational element 34a. It is contemplated that in some

embodiments of the present invention, a beveled cannula may be provided (such as in Fig. 5) and the rotational element may be somewhat recessed within the cannula, in that it does not extend further than a distal-most tip 80 thereof. 5 Thus, it is contemplated that as long as at least a portion of threading is exposed to tissue through the angled inlet, the tissue will be drawn into the inlet 42a and effectively removed upon rotation of the rotatable element.

Fig. 6 shows a cannula distal portion 40 similar to 10 that shown in Fig. 3. However the rotational element 34a is similar to that shown in Fig. 5, having narrow helical threading 56a, and a flat tip 53a rather than the rounded tip 53 shown in Fig. 3.

As shown in Figs. 1, 2 and 4, the apparatus 10 may 15 further comprise a collection chamber 70, for example, defined by a subhousing 72 removably engaged to the housing 22. More specifically, the collection chamber 72 is in fluid communication with a proximal portion 76 of the cannula 30. For example, the collection chamber 70 is 20 adapted to collect, temporarily contain, and allow analysis of breast tissue or other material, for example during and/or after the tissue removal procedure.

Generally, the collection chamber 70 is structured to 25 contain material that is drawn from the surgical site. The removed material enters the collection chamber 70 as shown by arrows 74 in Fig. 2. The collection chamber 70 is preferably adapted to allow observation of the tissue material during the procedure. For example, the subhousing 72 may be transparent. In addition, the collection chamber 30 70 is preferably structured to allow quantification or measurement of the tissue, for example, the subhousing 72 may be provided with suitable indices (not shown) showing milliliters (ml) of material collected therein. After the tissue removal procedure, the breast tissue may be biopsied 35 using conventional means. As shown, a proximal portion 78

of the rotatable element 34 is circumscribed by the collection chamber 70.

It is further contemplated that in many applications of the present invention, the cannula 30 may alternatively or additionally be used as a passageway for introducing medication and other agents into the target region of the breast before or after the tissue removal, if desirable.

It can be appreciated that the present apparatus is less invasive and more passive in comparison to other percutaneous breast tissue removal devices in the art. Despite its simplicity and passivity, the present device is designed to be highly effective in removing soft tissue, for example, cystic materials, fatty tissue, and muscle tissue or other materials that may be found in the breast. Because there is no external suction source or supplemental aspiration required to pull material into the cannula, it can further be appreciated that the apparatus is smaller, safer and requires less monitoring than devices that include a separate or external source of suction or additional idler shafts for removing material.

It is also to be appreciated that the apparatus of the present invention may be modified to include a connector for enabling the handpiece to be connected to an external aspiration source. In this case, means for monitoring the vacuum level in the cannula is preferably provided in order to indicate and prevent buildup of excess vacuum in the event the cannula becomes clogged for example.

While this invention has been described with respect to various specific examples and embodiments, it is to be understood that the invention is not limited thereto and that it can be variously practiced within the scope of the following claims.